L Number	Hits	Search Text	DB .	Time stamp
1	205	toxicant same (dissociat\$4 or inhibit\$4)	USPAT;	2004/08/24 12:09
		same (binding or bind or bound)	US-PGPUB;	
			EPO;	
			DERWENT	
2	2	toxicant same (dissociat\$4 or inhibit\$4)	USPAT;	2004/08/24 12:13
		same (binding or bind or bound) same	US-PGPUB;	
		immobili\$4	EPO;	
			DERWENT	
3	2	toxicant same (dissociat\$4 or inhibit\$4 or	USPAT;	2004/08/24 12:14
		reduc\$5 or prevent\$4) same (binding or	US-PGPUB;	
		bind or bound) same immobili\$4	EPO;	
	_		DERWENT	
4	. 7	toxicant same (dissociat\$4 or inhibit\$4 or	USPAT;	2004/08/24 12:14
		reduc\$5 or prevent\$4) same immobili\$4	US-PGPUB;	
			EPO;	
L			DERWENT	



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COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION

0.21

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FULL ESTIMATED COST

0.21 0

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=> toxicant and (dissociate or dissociation or inhibit or prevent or reduction or reduced or reduce) and (immobilized or immobilization or immobilizing)

L1 2 FILE AGRICOLA
L2 4 FILE BIOTECHNO
L3 0 FILE CONFSCI
L4 0 FILE HEALSAFE
L5 0 FILE IMSDRUGCONF

L6 3 FILE LIFESCI

L7 0 FILE MEDICONF

L8 1 FILE PASCAL

TOTAL FOR ALL FILES

10 TOXICANT AND (DISSOCIATE OR DISSOCIATION OR INHIBIT OR PREVENT OR REDUCTION OR REDUCED OR REDUCE) AND (IMMOBILIZED OR IMMOBILIZ ATION OR IMMOBILIZING)

=> dup rem

ENTER L# LIST OR (END):19

DUPLICATE IS NOT AVAILABLE IN 'IMSDRUGCONF, MEDICONF'. ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE

PROCESSING COMPLETED FOR L9

6 DUP REM L9 (4 DUPLICATES REMOVED)

=> d l10 ibib abs total

L10 ANSWER 1 OF 6 LIFESCI COPYRIGHT 2004 CSA on STN

8/24/J

2002:53344 LIFESCI ACCESSION NUMBER:

Embryonic development assay with Daphnia magna: application TITLE:

to toxicity of aniline derivatives

Abe, Tatsuo; Saito, Hotaka*; Niikura, Yoshiyuki; Shigeoka, AUTHOR:

Tadayoshi; Nakano, Yoshio

Department of Environmental Chemistry and Engineering, CORPORATE SOURCE:

Tokyo Institute of Technology, 4259 Nagatsuta-cho,

Midori-ku, Yokohama 226-8502, Japan; E-mail:

gfs@ankaken.co.jp

Chemosphere, (20011100) vol. 45, no. 4-5, pp. 487-495. SOURCE:

ISSN: 0045-6535.

Journal DOCUMENT TYPE:

FILE SEGMENT:

Х

LANGUAGE:

English

SUMMARY LANGUAGE:

English

An assay system using Daphnia magna embryos was applied to investigate the adverse effects of aniline derivatives. The data were compared with our previous data for chlorophenols. This new assay provides useful information to evaluate the toxicity of chemicals and the differences in sensitivity between the life stages. The effects of 15 aniline derivatives on embryonic development of D. magna embryos were determined. At the start of exposure, 2-6-h old eggs (between stages 1 and 2, round in shape, diameter approx. 400 mu m), were used. In control and solvent control groups, embryonic development from an egg to a free-swimming animal proceeded completely within 3 days with more than 90% hatchability. Median effective concentrations (EC sub(50)s) to reduce the numbers hatched were determined and gross morphological abnormalities of hatched animals recorded. Anilines induced no obvious morphological abnormalities and no developmental delay although premature deaths occurred. However, they affected the number of embryos hatching in a dose-dependent manner. In addition, this embryo assay was more sensitive to aniline derivatives (except for aniline) than acute juveniles immobilization assay. Ratios of 48-h EC sub(50) (juvenile)/3-day EC sub(50) (embryo) for eight anilines were greater than 5.0. Particularly, the ratios of 4-methyl-, 4-ethyl- and 3-methylaniline were 77, 23 and 11, respectively. EC sub(50)s for embryos and juveniles were poorly correlated (r = 0.41). This indicated that the sensitivities of the two life stages were different to the effects of anilines. EC sub(50)s were poorly correlated (r = -0.097) with the log K sub(ow) (1-octanol/water partition coefficient). These results were compared with previous results for phenols.

ANSWER 2 OF 6 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2004) on STN DUPLICATE 1

ACCESSION NUMBER:

2000:65945 AGRICOLA

DOCUMENT NUMBER:

IND22059087

TITLE:

In vitro phytotoxicity screening test using

immobilized spinach thylakoids.

AUTHOR(S):

Laberge, D.; Chartrand, J.; Rouillon, R.; Carpentier,

R.

AVAILABILITY:

DNAL (OH545.A1E58)

SOURCE:

Environmental toxicology and chemistry, Dec 1999. Vol.

18, No. 12. p. 2851-2858

Publisher: Pensacola, Fla. : SETAC Press.

CODEN: ETOCDK; ISSN: 0730-7268

NOTE:

Includes references Florida; United States

PUB. COUNTRY: DOCUMENT TYPE:

Article

U.S. Imprints not USDA, Experiment or Extension

FILE SEGMENT:

LANGUAGE: English Several pollutants found in water inhibit the photosynthetic

electron transport chain, and therefore affect the growth of phytoplankton and aquatic plants. In this study, thylakoid membranes isolated from

spinach leaves were used in a microelectrochemical cell to generate photocurrent. The toxic effect of an inhibitor is observed by a decrease in the photocurrent. To improve the stability of their biological functions, the thylakoid membranes were immobilized in an albumin-glutaraldehyde cross-linked matrix. The developmental work of this phytotoxicity test was done by using the herbicide atrazine as the reference toxicant. Results on reproducibility were in the range generally accepted for standardized bioassays. The phytotoxicity of herbicides from various chemical classes including photosynthetic and nonphotosynthetic inhibitors was evaluated. Toxicity responses of the immobilized thylakoid test to photosynthetic inhibitors compared favorably with literature data for the algal growth inhibition test using Selenastrum capricornutum. The detection capabilities of the photosynthetic microassay for cyanazine, metribuzin, diuron, and propanil met the recommandation for the water quality guidelines for raw water. Characteristics of this in vitro approach such as rapidity, experimental simplicity, and cost effectiveness are also discussed.

L10 ANSWER 3 OF 6 BIOTECHNO COPYRIGHT 2004 Elsevier Science B.V. on STN

ACCESSION NUMBER:

1997:28326014 BIOTECHNO

TITLE:

Effects of immobilization restraint on

syrian golden hamsters

AUTHOR:

King-Herbert A.P.; Hesterburg T.W.; Thevenaz P.P.; Hamm T.E. Jr.; Moss O.R.; Janszen D.B.; Everitt J.I.

CORPORATE SOURCE:

Dr. A.P. King-Herbert, CIIT, P.O. Box 12137, Research

Triangle Park, NC 27709, United States.

SOURCE:

Laboratory Animal Science, (1997), 47/4 (362-366), 14

reference(s)

CODEN: LBASAE ISSN: 0023-6764

DOCUMENT TYPE:

Journal; Article

COUNTRY:

AΒ

United States

LANGUAGE:

English

SUMMARY LANGUAGE:

English

AN 1997:28326014

4 BIOTECHNO

Rodent nose-only inhalation toxicology systems comprise whole-body immobilization in plastic restraint tubes. This method of restraint is known to have a variety of effects on animals. In the studies reported here, two independent toxicology laboratories examined the effects of inhalation tube restraint in Syrian golden hamsters, a species that has recently gained importance in inhalation studies of fibrous particulates. Body weight, food and water consumption, core body temperature, and plasma cortisol and corticosterone concentrations were assessed in animals immobilized in nose- only inhalation tubes, and the results were compared with those from unrestrained cage-control animals. Animals were immobilized for either 6 h/ day, 5 days/week for 13 weeks (subchronic), or 4 h/day for 14 consecutive days (subacute), mimicking exposure conditions commonly used in nose-only inhalation studies. Tube restraint was found to induce a marked decrease in body weight, which increased in response to cessation of restraint. The body weight decrement was associated with significant differences in food and water consumption between the restrained and control groups in the subacute study and only food consumption in the subchronic study. During the restraint period, core body temperature in the immobilized animals increased slightly but not above the normal range for this species. Plasma cortisol and corticosterone concentrations were not significantly increased with use of restraint, compared with values in controls. Immobilization-associated body weight depression in Syrian golden hamsters is important for the evaluation of nose-only inhalation study results because many normal physiologic parameters, as well as toxicant-induced effects, are associated with body weight status.

L10 ANSWER 4 OF 6 LIFESCI COPYRIGHT 2004 CSA on STN ACCESSION NUMBER: 97:97582 LIFESCI

Ecotoxicity assessment of the aquatic environment around TITLE:

Lake Kojima, Japan

Okamura, H.; Luo, R.; Aoyama, I.; Liu, D. AUTHOR:

Res. Inst. for Bioresources, Okayama Univ., 2-20-1 Chuo, CORPORATE SOURCE:

Kurashiki, Okayama 710, Japan

ENVIRON. TOXICOL. WATER QUAL., (1996) vol. 11, no. 3, pp. SOURCE:

213-221.

Meeting Info.: 7. International Symposium on Toxicity

Assessment. [np]. ISSN: 1053-4725.

DOCUMENT TYPE: TREATMENT CODE: Journal Conference

FILE SEGMENT:

X; K English

LANGUAGE: SUMMARY LANGUAGE: English

To reduce the impact of chemical substances on the aquatic ecosystem, it is essential to understand their ecotoxicological properties in the natural aquatic environment. Consequently, we conducted an ecotoxicological study on the aquatic environment around Lake Kojima, a man-made lake located in the southwest of Japan. Lake Kojima receives its chemical inputs mainly from two rivers that flow through various agricultural and industrial areas. For ecotoxicity screening, surface water and sediment samples were collected 4 times in 1993 from 16 preselected sites. Then, the solutes in the filtered surface water were concentrated by ODS resin, and the organic chemicals in the suspended solids (SS) and sediments were extracted by acetone. A battery of five ecotoxicity tests (agar plate test using bacteria and yeast, algal growth inhibition test. Daphnia magna immobilization test, and root elongation test using lettuce seeds) was used to assess these extracts. The results show that the surface water extracts had a lethal effect on D. magna, the SS extracts suppressed algal growth, and the sediment extracts were inhibitory to the growth of yeast. A significant inhibitory effect by the sediment extracts from 4 lake sites and 3 river sites was detected by these ecotoxicity tests. Attempts also were made to identify the putative ecotoxic chemicals in the collected samples. Elementary sulfur was identified as one of the major toxicants in the sediment extracts that were inhibitory to the yeast growth. Moreover, samples of surface water around Lake Kojima, collected weekly from June to September in 1994, were found to contain three pesticides and were toxic to D. magna. But the concentration of the pesticides detected was too low to cause Daphnia immobilization. It is believed that the toxicity of the water extracts was mainly due to the combined toxic effect of natural and man-made components.

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ACCESSION NUMBER:

94:9318 AGRICOLA

DOCUMENT NUMBER:

IND20367004

TITLE:

Toxicity of tributyltin chloride to anaerobic nitrogen

transformations in sediment and porewater.

AUTHOR (S):

Bergeron, V.; Blais, J.S.; Wharf, I.; Marshall, W.D.

AVAILABILITY: DNAL (QH540.J6)

SOURCE:

Journal of environmental quality, July/Sept 1993. Vol.

22, No. 3. p. 528-536

Publisher: Madison: American Society Of Agronomy,

CODEN: JEVQAA; ISSN: 0047-2425

NOTE:

Paper presented at the USDA-ARS Beltsville Agricultural Research Center Symposium XVII, "Agricultural Water Quality Priorities, A Team Approach to Conserving Natural Resources, " May 4-8,

1992, Beltsville, MD. Includes references

PUB. COUNTRY:

United States; Wisconsin

DOCUMENT TYPE:

Article

FILE SEGMENT:

U.S. Imprints not USDA, Experiment or Extension

English

LANGUAGE: The influence of tributyltin chloride (TBTCI) on N transformations in anoxic sediment cultures, during approximately 14-d incubations, was studied using acetylene inhibition and acetylene reduction techniques as measures of microbially mediated denitrification and dinitrogen fixation, respectively. The accumulation of N2O, CO2 and C2H4 with time was modeled with a best-fit polynomial to detect statistically significant differences between treatments and with a three-segment continuous line model to assess lag times, rates of accumulation and rates of subsequent loss of these gases from the headspace. In sediment cultures, the presence of up to 100 mg L-1 of TBT had a barely detectable influence on these transformation. However, the analogous processes in porewater, prepared by centrifuging sediment slurry at 3 000 X g for 30 min, were appreciably modified by the presence of > 1 mg L-1 of this toxicant. Although the two media were different in terms of their denitrifying potential and their fermenting capacity, dose-related responses in the porewater were evident for both processes. Apparently the presence of particulate matter in the sediment slurry appreciably attenuated the inhibitory effects of the toxicant. Moreover, a portion of the denitrifiers in cultures of porewater developed a resistance to 100 mg L-1 of TBT in the medium and reduced added nitrate stoichiometrically. When transferred to autoclaved medium containing the same level of toxicant, aliquots of the resistant culture stoichiometrically reduced the added nitrate after a shorter lag time. The resistance was not the result of metabolic detoxification as indicated by the recovery of 74% of toxicant in an unchanged form, after 15 d incubation, using analytical methods which would have detected a 1% conversion of TBT to either Bu2Sn2+ or to BuSn3+. When added, at 10 mg L-1, to autoclaved porewater, an appreciable portion of the toxicant became associated with residual particulate matter (the fraction removed by centrifugation at 10 000 X g but not by 3 000 X g) with only approximately 9% remaining in the supernatant fluid. When 0.5 mL of TBT-resistant culture was added to the identical matrix and incubated, the TBT in the supernatant phase was reduced below the limit of detection and only 40% was recovered in the particulate/microbial cell fraction. Thus, both particulate materials and microbial growth immobilized TBT serving to limit its concentration in the surrounding water.

ANSWER 6 OF 6 BIOTECHNO COPYRIGHT 2004 Elsevier Science B.V. on STN L10 DUPLICATE

ACCESSION NUMBER:

BIOTECHNO 1991:21345696

TITLE:

Immobilized microbe bioreactors for waste

water treatment

AUTHOR:

Portier R.J.; Miller G.P.

CORPORATE SOURCE:

Aquatic/Industrial Toxicology Laboratories, Institute for Environmental Studies, Louisiana State University,

Baton Rouge, LA 70803, United States.

SOURCE:

Waste Management and Research, (1991), 9/5 (445-451)

CODEN: WMARD8 ISSN: 0734-242X

DOCUMENT TYPE:

Journal; Conference Article

COUNTRY: LANGUAGE: United Kingdom

English English

SUMMARY LANGUAGE: 1991:21345696 **BIOTECHNO** AN

The application of adapted microbial populations immobilized on AB a porous diatomaceous earth carrier to pre-treat and reduce toxic concentration of volatile organics, pesticides, petroleum aliphatics and aromatics has been demonstrated for several industrial sites. In the pre-treatment of industrial effluents and contaminated ground-waters, these bioreactors have been used to optimize and

reduce the cost of conventional treatment systems, i.e. steam stripping, carbon adsorption and traditional biotreatment. Additionally, these systems have been employed as seeding devices for larger biotreatment systems. The cost effective utilization of an immobilized microbe reactor system for water supply regeneration in a microgravity environment is presented. The feasibility of using immobilized biomass reactors as an effluent treatment technology for the biotransformation and biodegradation of phenols, chlorinated halocarbons, residual oils and lubricants was evaluated. Primary biotransformation tests of two benchmark toxicants, phenol and ethylene dichloride at concentrations expected in life support effluents were conducted. Biocatalyst supports were evaluated for colonization potential, surface and structural integrity, and performance in continuous flow bioreactors. The implementation of such approaches in space will be outlined and specific areas for interfacing with other non-biological treatment approaches will be considered for advanced life support, tertiary waste water biotreatment.